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EVALUATION OF CANDIDATE WORKING FLUID FORMULATIONS FOR THE ELECTROTHERMAL - CHEMICAL WIND TUNNEL

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Introduction

A new hypersonic test facility which can simulate conditions typical of atmospheric flight at Mach numbers up to 20 is currently under study at the NASA/LaRC Hypersonic Propulsion Branch. In the proposed research it was suggested that a combustion augmented electrothermal wind tunnel concept may be applied to the planned hypersonic testing facility. The purpose of the current investigation is to evaluate some candidate working fluid formulations which may be used in the chemical-electrothermal wind tunnel.

Work Done

The efforts in the initial phase of this research were concentrated on acquiring the code used by GASL to model the electrothermal wind tunnel and testing it using the conditions of GASL simulation. The early version of the general chemical kinetics code (GCKP84) was obtained from NASA and the latest updated version of the code (LSENS) was obtained from the author Dr. Bittker. Both codes are installed on a personal computer with a 486 25 MHz processor and 16 Mbyte RAM. Since the available memory was not sufficient to debug LSENS, for the current work GCKP84 was used.

The Effect of NO₂ Reactions: As a first step we tried to reproduce the results obtained by GASL for air-argon mixtures. For this purpose the GASL kinetic and thermodynamic data provided in reference (1) are used. Although kinetic constants for reactions involving NO₂ (reactions 6 and 7 in the GASL table) were given in Table 7 of reference (1), it was indicated in the report that these reactions were not considered in the computations. Private communication with O. F. Rizkalla confirmed this and the reason given for the exclusion of these reactions were the unavailability of high temperature thermodynamic data and the expectation that their influence on the test gas composition will be negligible.

We have obtained high temperature thermodynamic data for NO2

and several other species of importance to our analysis². The data obtained contains two sets of coefficients for the ranges 200 to 1000 K and 1000 to 10,000 K. For some species the upper limit is 20,000 K. Since GASL has used three temperature interval fit for their thermodynamic data, the NO₂ data could not be incorporated to their thermodynamic data, and the McBride data was used in our studies as a complete set.

The initial conditions for the reactive nozzle flow were taken from Table 6 of reference (1). A 5° nozzle with the specified throat area was used. The nozzle area, A in cm², is given by

$$A = 0.0240466 x^2 + 0.198988 x + 0.41166$$

In our simulations the reactions and accompanying rate data given in Table 7 of the GASL study¹ are used. For some runs the kinetic data for reaction 6 was replaced by the data from a test case supplied by Dr. Bittker with LSENS³, and the results shown in the second row were obtained using the high temperature kinetic data for reactions 6 and 7 reported by Jachimowski⁴. The result of our simulations are compared with the GASL results¹ in Table 1.

Considering the fact that the GASL results are read from a graph, the NO, O, and O_2 mole fractions have been reproduced by using both the GASL thermodynamic data and the McBride data. In fact, McBride Thermodynamic data appears to produce results closer to those of GASL. The only major discrepancy between our simulations and the GASL results is the nozzle length required to produce an exit Mach number of 16. The nozzle length from Figure 16 of reference 1 is about 8.9 ft while our simulations indicate a nozzle length of about 8.7 m.

The first 6 rows of the table show the effect of NO_2 kinetics on the composition of the gas in the nozzle. The main effect of the NO_2 reactions is on the oxygen atom concentration. When these reactions are excluded from the calculations the oxygen atom mole fraction at the nozzle exit is 1.24E-2 while with the NO_2 reactions the oxygen atom mole fraction falls below 7.25E-5 for all rate

data. The effect of NO_2 reactions on the NO mole fraction is not that significant. The NO mole fraction is 4.84E-2 when the reactions are excluded and about 5.6E-2 when they are included.

Table 1. Effect of NO₂ reactions on the test gas composition with air as the working fluid.

Reaction	Conditions at 800 cm (Y; in mole frac.)					
Information	Т (К)	Y _{NO}	Yo	У ₀₂	M _a	
NO ₂ Reactions Included GASL Reaction Data	271.66	5.592E-2	5.625E-7	0.1813	15.367	
NO ₂ Reactions Included Data from Jachimowski	270.90	5.604E-2	7.253E-5	0.1813	15.388	
NO ₂ Reactions Included Bittker's Data for 6	267.04	5.648E-2	6.883E-14	0.1811	15.477	
Reaction 7 Excluded GASL Data for 6	322.81	3.653E-2	2.715E-3	0.1782	14.131	
Reaction 7 Excluded Bittker's Data for 6	291.57	3.352E-2	1.387E-3	0.1777	14.910	
NO ₂ Reactions Excluded McBride Thermo Data	263.20	4.838E-2	1.246E-2	0.1778	15.505	
NO ₂ Reactions Excluded GASL Thermo Data	260.8	5.106E-2	1.067E-2	0.1776	15.55	
GASL Results at Nozzle Exit		4.75 E-2	1.9 E-2	0.173	16.0	
NO ₂ Reactions excluded McBride Data-Exit Cond.	247.44	4.838E-2	1.244E-2	0.1778	16.00	

Mach 16 free jet conditions for air, air containing 10% ammonia, air containing water equivalent to the hydrogen in 10% ammonia mixture, and a mixture obtained by replacing some nitrogen in air by ammonia to give 10% ammonia. Computations for air and air + 10% NH₃ mixtures were done for 5 and 10 degree nozzles while for the others only a 5 degree nozzle was used. The total enthalpy was

3159.5 cal/g and the mass flow rate was 13302 g/s for all computations. The specified total enthalpy is assumed to be provided by electrical energy input to the feed gas mixture. The mixing chamber pressure was 4690 atm. A lower pressure, 2554 atm, was also tried for air and air + 10% NH₃ mixture. These values were chosen to keep the conditions used in this study similar to the conditions of the GASL study. In all computations reactions involving NO₂ were included and the rate data reported by Jachimowski⁽⁴⁾ were used for these reactions. For all the other reactions GASL kinetics⁽¹⁾ were employed.

The computation procedure was as follows:

- Find the initial mixture temperature to give the desired total enthalpy in the mixing chamber. Chemical and thermal equilibrium was assumed to exist in the mixing chamber. The equilibrium conditions and gas composition were found using the equilibrium combustion option of the GCKP.
- Pressure and temperature at the throat are calculated from the reservoir conditions found in the first step using

$$p^* = p_0 \left(\frac{2}{\gamma + 1} \right)^{\frac{\gamma}{\gamma - 1}}$$
 $T^* = T_0 \left(\frac{2}{\gamma + 1} \right)$

Gas composition at the throat was assumed to be the same as that in the mixing chamber.

- The gas velocity was adjusted to give an initial Mach

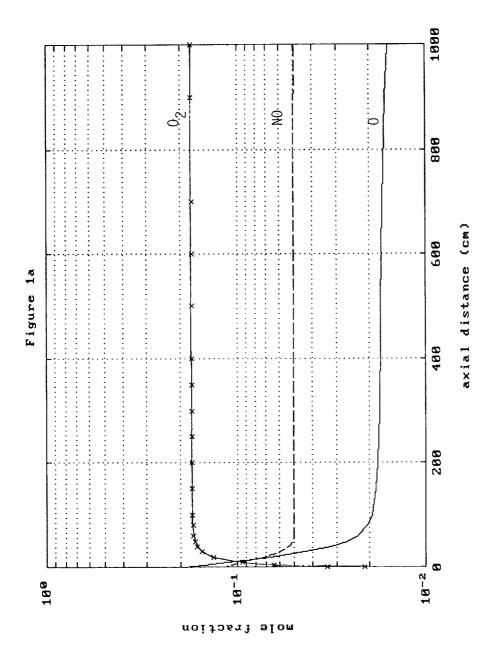
number slightly above 1 (usually around 1.025). In some cases, due to the reactions taking place, just after the throat, temperature increases and Mach number decreases and the flow is choked not at the throat but at some distance from it in the expanding section of the nozzle. This necessitates starting Mach numbers even as high as 1.16 at the throat.

 Adjust the throat area to give the desired mass flow rate.

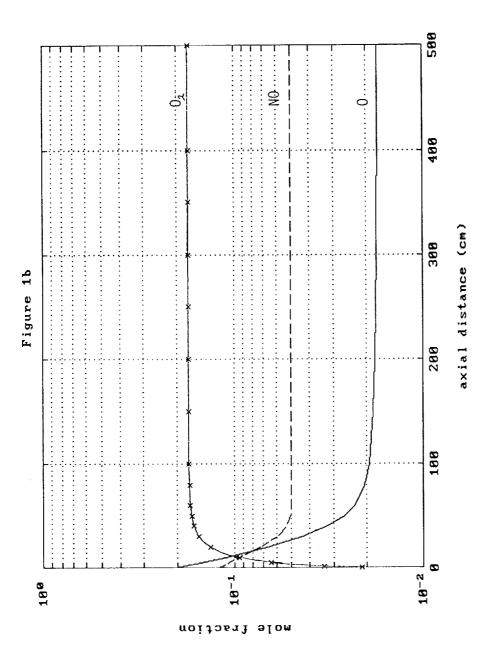
Results: The results are shown in Figures 1 through 9. In each figure the first chart shows the variation of gas composition up to an axial distance of 1000 cm while the second chart covers the region up to an axial distance of 500 cm and is a blown up version of the first one. The results indicate that for the conditions specified, the flow in the nozzle is essentially frozen beyond 500 cm and for this reason the flow conditions and gas compositions at 500 cm are summarized in Table 2.

The conclusions derived from Table 2 can be summarized as follows:

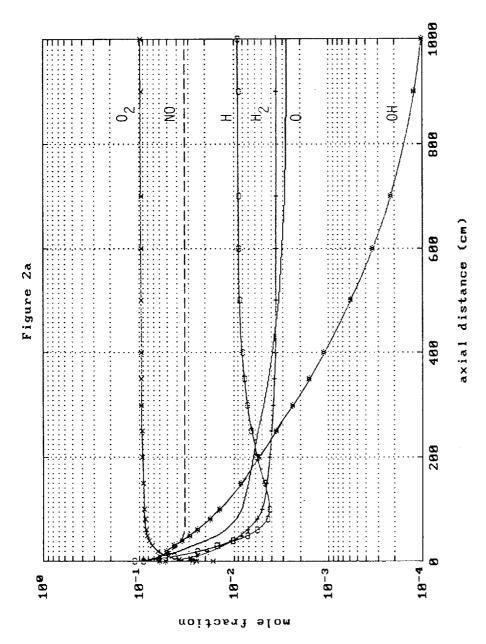
- The addition of $\mathrm{NH_3}$ or $\mathrm{H_2O}$ to air decreases the Mach number at the nozzle exit. This is mainly due to smaller γ values of these mixtures. To achieve Mach 16 at the nozzle exit the expansion angle of the nozzle after about 300 cm from the throat should be increased gradually. An initial section with 5 degree expansion is necessary to preserve the high rate of



Gas composition during reactive flow through 5^o nozzle. Mixing chamber pressure 2554 atm. Air feed

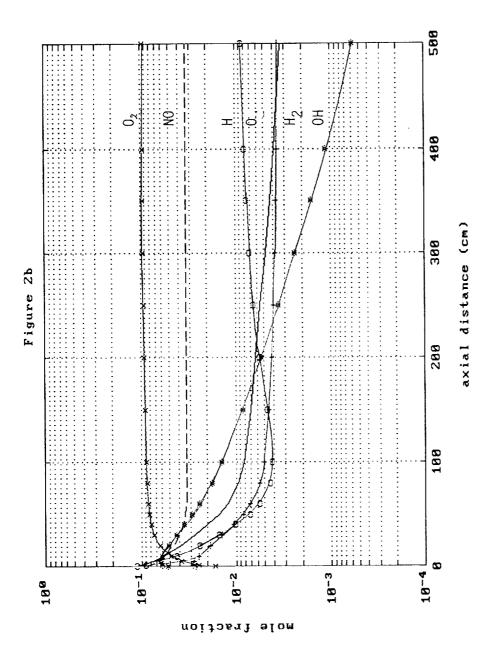


Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing chamber pressure 2254 atm.

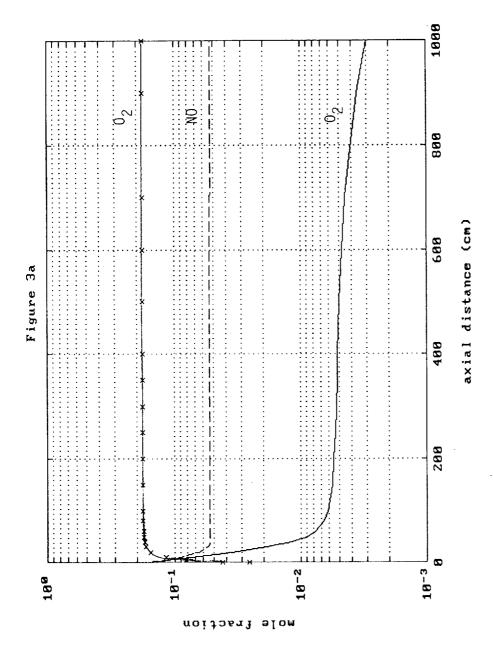


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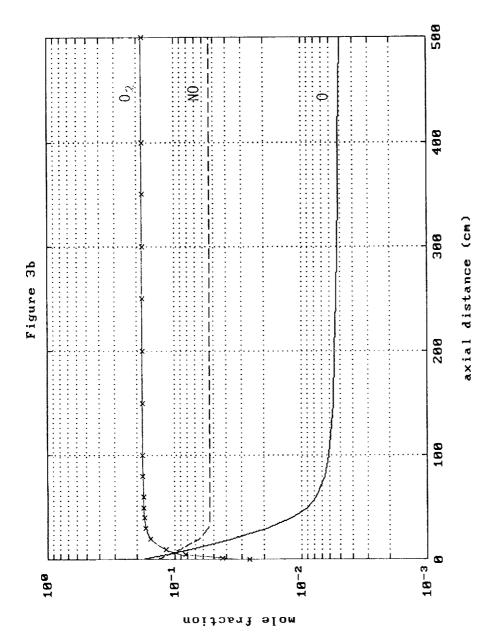
Gas composition during reactive flow through $\mathbf{5}^{\rm O}$ nozzle. Mixing chamber pressure 2554 atm. Feed is $10\%~{\rm NH}_3$ in air.



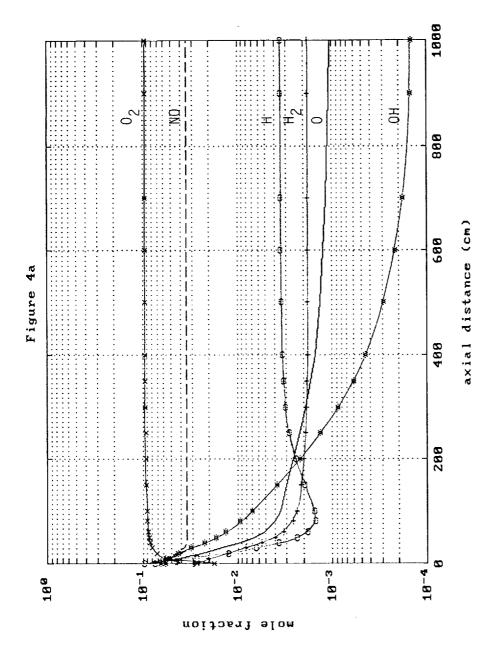
Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing chamber pressure 2554 atm. Feed is $10\%~{
m NH}_3$ in air.



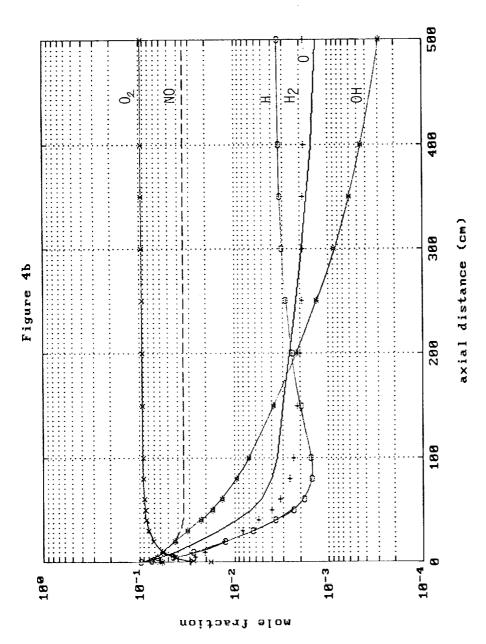
Gas composition during reactive flow through 5° nozzle. Mixing chamber pressure 4690 atm. Air feed.



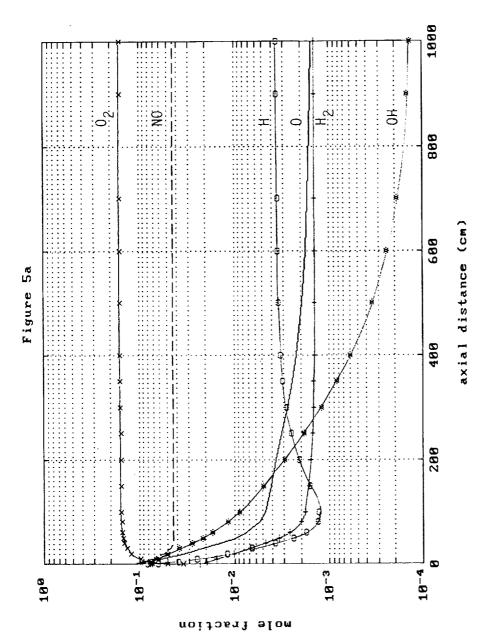
Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing chamber pressure $4690~{\rm atm.}$



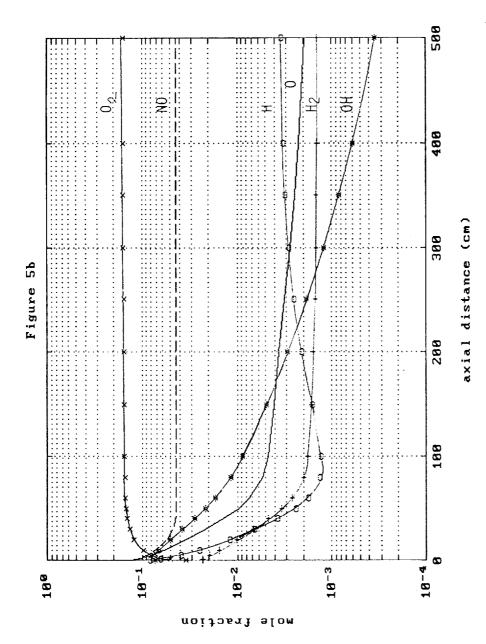
Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing chamber pressure 4690 atm. Feed is $10\%~{\rm NH}_3$ in air.



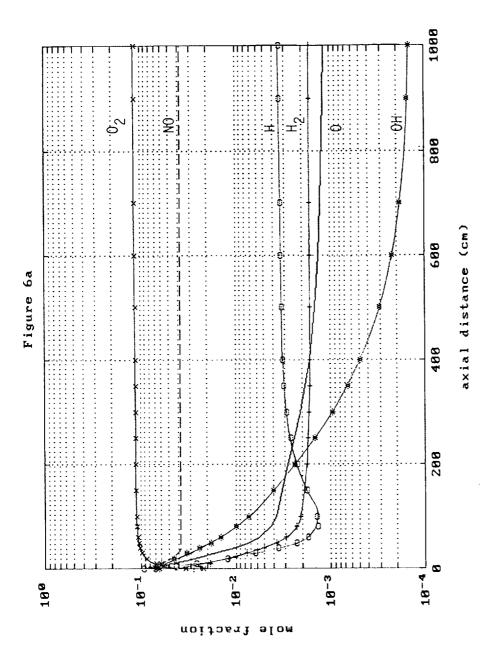
Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing chamber pressure 4690 atm. Feed is $10\%~{\rm NH}_3$ in air.



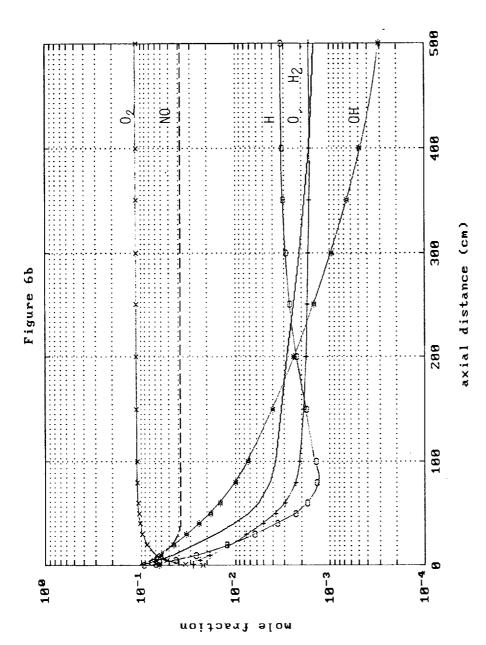
chamber pressure 4690 atm. Feed is air + $\mathrm{H}_2\mathrm{O}$ equivalent to H in Gas composition during reactive flow through 5^{0} nozzle. Mixing 10% NH₃.



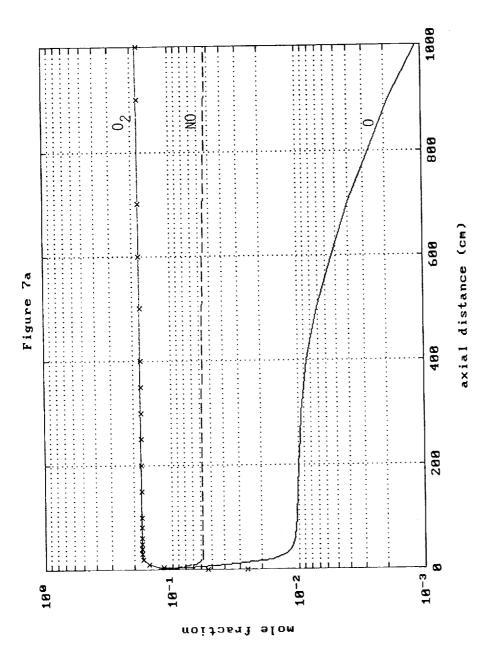
chamber pressure 4690 atm. Feed is air + $\mathrm{H}_2\mathrm{O}$ equivalent to H in Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing



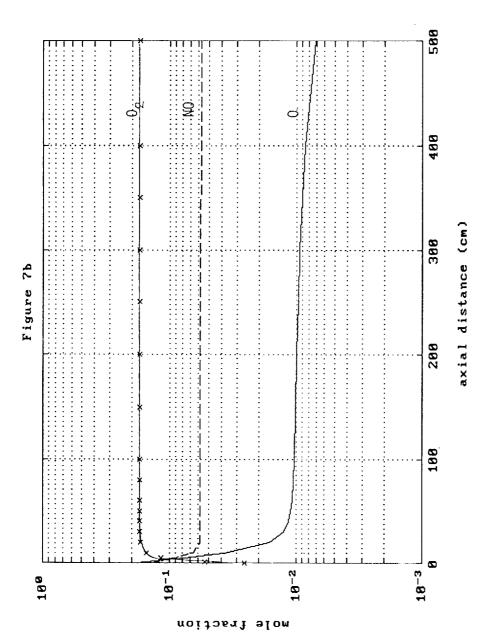
Gas composition during reactive flow through $5^{\rm O}$ nozzle. Mixing chamber pressure 4690 atm. Some $\rm N_2$ in air replaced to give 10% NH $_3$ in the feed.



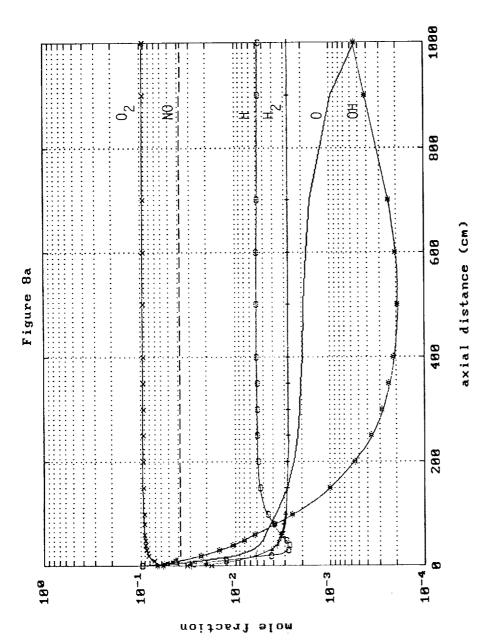
chamber pressure 4690 atm. Some $\rm N_2$ in air replaced to give $10\%~\rm NH_3$ Gas composition during reactive flow through 5^{0} nozzle. Mixing in the feed.



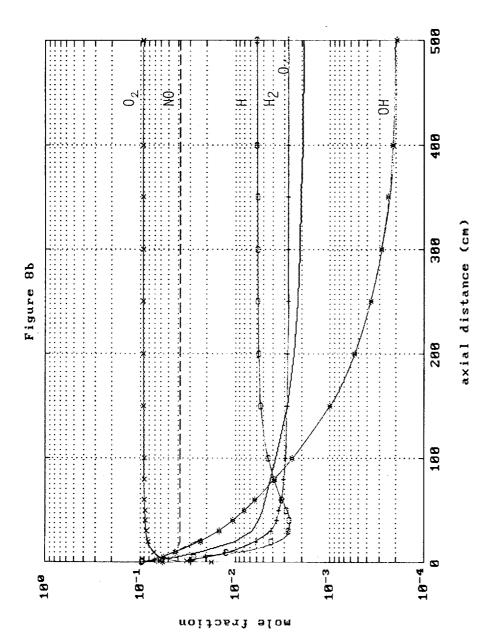
Gas composition during reactive flow through $10^{\rm O}$ nozzle. Mixing chamber pressure 4690 atm. Air feed.



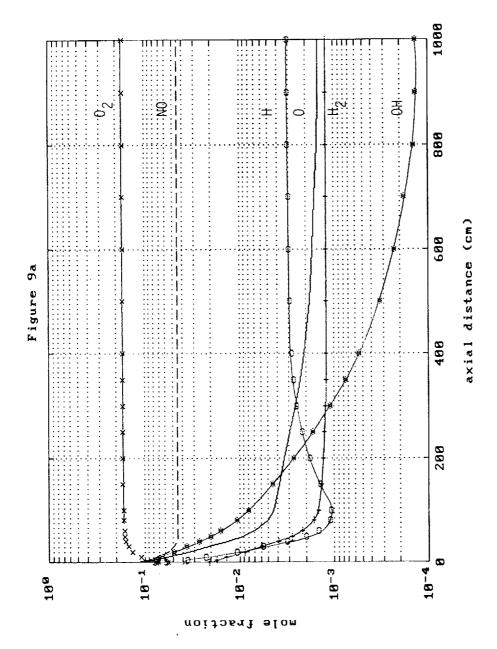
Gas composition during reactive flow through EV nozzle. Mixing chamber pressure 4690 atm.



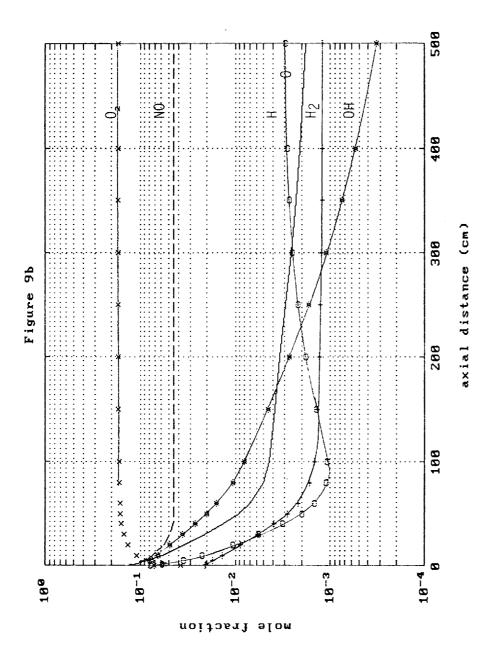
Gas composition during reactive flow through $10^{\rm O}$ nozzle. Mixing chamber pressure $4690 \ \mathrm{atm.}$ Feed is $10\% \ \mathrm{NH_3}$ in air.



Gas composition during reactive flow through $10^{\rm O}$ nozzle. Mixing chamber pressure 4690 atm. Feed is $10\%~\mathrm{NH}_3$ in air.



chamber pressure 4690 atm. Feed is $10\%~\mathrm{NH_3}$ in oxygen-enriched air. Gas composition during reactive flow through 5° nozzle. Mixing



chamber pressure 4690 atm. Feed is $10\%~\mathrm{NH_3}$ in oxygen-enriched air. Gas composition during reactive flow through 50 nozzle. Mixing

recombination reactions.

Table 2. Summary of results for reactive nozzle flows for various gas compositions. Ma = Mach Number, y_i = mole fraction of species i.

Gas composi- tion	Air		Air + 10% NH ₃			Some N ₂ in air replaced by NH ₃ to give 10% NH ₃	air + H ₂ O equivalent to H in 10% NH ₃	
P ₀ (atm)	<u>2254</u>	<u>4690</u>	<u>4690</u>	2254	4690	4690	4690	4690
θ (deg .)	5	5	10	5	5	10	5	5
Conditions at 500 cm from nozzle throat								
P (atm)	.0076	.0062	.0009	.0107	.0087	.0012	.0086	.0091
T (K)	636.5	517.6	320.8	863.9	703.4	409.7	714.2	749.1
Ma	10.18	11.37	14.45	8.89	9.91	12.95	9.85	9.68
γ	1.375	1.386	1.399	1.337	1.353	1.384	1.352	1.346
y _o	.0174	.0050	.0070	.0033	.0013	.0019	.0014	.0019
Умо	.0500	.0535	.0570	.0333	.0347	.0381	.0376	.0434
y _{o2}	.1740	.1797	.1753	.0953	.0942	.0932	.1129	.1578
y _{ii}				.0086	.0034	.0059	.0033	.0034
Уон				5.9E-4	2.9E-4	2.0E-4	3.0E-4	3.5E-4
y _{H2}				.0036	.0018	.0028	.0016	.0014
y _{H20}				.1363	.1416	. 1357	.1419	.1385
y ₀ /y ₀₂	.1000	.0278	.0399	.0346	.0138	.0204	.0124	.0120
Y _{NO} /Y _{OZ}	.2874	.2977	.3252	.3494	.3684	.4088	.3330	.2750

- As expected, a larger nozzle expansion angle freezes the reactions earlier. As a consequence, the mole fractions of O, NO, and H are larger while those of O₂ and H₂ are smaller when the flow is expanded faster. The OH mole fraction is smaller for 10 degree expansion. The most significant changes are observed for the O mole fraction in pure air and H mole fraction in air containing 10% NH₃.
- When column 2 is compared to column 5 it is seen that an original gas mixture containing 10% $\rm NH_3$ results in a test gas

with significantly lower O and NO mole fractions at the expense of having some H and OH in the test gas. The H mole fraction is over an order of magnitude smaller than that of NO and the OH mole fraction is over an order of magnitude smaller than that of H. Since the O_2 mole fractions are different, to obtain a better comparison, O/O_2 and NO/O_2 ratios are given in rows 15 and 16, respectively. These entries indicate that although the addition of ammonia to air is effective in reducing the O/O_2 ratio, it increases the NO/O_2 ratio due to the presence of a larger number of N atoms.

- Since some air in the feed gas is replaced by NH, in the 10% ammonia containing feed, the O_2 mole fraction in the test gas for this case is much lower and the number of N atoms present is much higher than with pure air feed. To obtain a better comparison, the results for a feed gas in which some nitrogen is replaced by ammonia to give 10% $\mathrm{NH_{3}}$, are given in column 7. Even in this case the O_2 mole fraction is lower than with pure air feed because some oxygen is used to form water. The 0/02 ratio is lower than that obtained with pure air feed and the NO/O2 ratio is only slightly higher. There was a significant improvement over the 10% $\mathrm{NH_3}$ feed. The H and $\mathrm{H_2}$ mole fractions are also lower with only insignificant increases in OH and Since the OH concentration continues to decrease up to the nozzle exit unlike the other species whose concentrations are frozen after about 500 cm after the nozzle throat, it is expected that the OH mole fraction in the test gas will be lower than the value reported in Table 2.
- To see if the beneficial effect of NH_3 is due to the H_2O formed, the results obtained for a gas feed containing H_2O equivalent to H in the 10% NH_3 feed are included in column 8 of Table 2. These results show that the feed containing H_2O gives the best O/O_2 and NO/O_2 ratios. Even the H_2 and H_2O mole fractions are lower than those in column 6 with only slight increases in H and OH.

Table 3. Summary of results for reactive nozzle flows for various gas compositions. Mixing chamber pressure 4690 atm, nozzle expansion 5 degrees. Ma = Mach number, y_i = mole fraction of species i.

Gas composi- tion	Air	Some N ₂ in air replaced by NH ₃ to give 10% NH ₃ + .08 mol O ₂ per mol	Air + H ₂ O equivalent to H in 10% NH ₃				
P ₀ (atm)	<u>4690</u>	4690	4690				
θ (deg .)	5	5	5				
Conditions at 500 cm from nozzle throat							
P (atm)	.00618	.00891	.00911				
T (K)	517.59	732.05	749.08				
Ma	11.37	9.759	9.678				
Υ	1.3856	1.3483	1.3462				
y _o	.004958	.001811	.001924				
Уно	.053517	.045193	.043374				
Yoz	.179664	.171333	.157830				
Ун		.002975	.003408				
Уон		.000324	.000354				
Ун2		.001204	.001430				
У _{н20}		.132001	. 138533				
y ₀ /y ₀₂	.027596	.010570	.012190				
y _{NO} /y _{O2}	. 297873	.263773	.274815				

- To compare the results for feed gas compositions giving similar O₂ mole fractions in the test gas, results were also obtained for a feed gas containing 10% NH₃ and enriched in oxygen to give an O₂ mole fraction similar to that in the test gas obtained with pure air. These results are presented in column 2 of Table 3 and compared with some of the results from Table 2. These results indicate that better O/O₂ and NO/O₂ ratios can be obtained with oxygen enriched air containing 10% NH₃ than air containing water in an amount containing same number of H atoms as in the 10% NH₃ mixture. In addition, the

mole fractions of H, OH, $\rm H_2$, and $\rm H_2O$ are all lower with the 10% NH $_3$ containing enriched air.

Conclusions

The results shown above indicate that the addition of nitrogen and hydrogen-containing compounds to oxygen-enriched air can improve the $0/0_2$ and $N0/0_2$ ratios and can have lower H, OH, H₂, and H₂O mole fractions than with feed gases containing H₂O. It appears that the liquid propellant HAN $(N_2H_4O_4)$ may even be a better additive than NH_3 due to its lower N/H ratio and high oxygen content. Other propellants such as HMX, RMX, and HNTO will produce some CO_2 which will reduce the specific heat ratio (thus improving the required stagnation conditions) and increase the rates of recombination reactions in the nozzle. The presence of CO_2 in the during combustion tests.

References

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